Mayer-Rokitansky-Küster-Hauser syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mayer-Rokitansky-Küster-Hauser syndrome. ORPHA:3109

Mayer-Rokitansky-Küster-Hauser (MRKH) syndrome describes a spectrum of Mullerian duct anomalies characterized by congenital aplasia of the uterus and upper 2/3 of the vagina in otherwise phenotypically normal females. It can be classified as either MRKH syndrome type 1 (corresponding to isolated utero-vaginal aplasia) or MRKH syndrome type 2 (utero-vaginal aplasia associated with other malformations) (see these terms).